Amendments to the Claims:

Please amend claims 1, 2 and 13, cancel claims 4-12 and 15-50 and add new claims 51-62. This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

- 1. (currently amended) A composition comprising an isolated nucleic acid molecule which encodes a Pvs25 polypeptide <u>having at least 95% identity to SEQ ID NO:4</u>, wherein the polypeptide induces an immune response in a susceptible organism that blocks the <u>transmission of malaria</u> and hybridizes under stringent conditions to SEQ ID NO:3.
- 2. (currently amended) <u>A The composition comprising an of claim 1,</u> wherein the isolated nucleic acid <u>having</u> has a sequence as shown in SEQ ID NO:3.
- 3. (original) A composition comprising an isolated nucleic acid molecule which encodes a Pvs25 polypeptide having an amino acid sequence as shown in SEQ ID NO:4.
 - 4-12 (canceled)
- 13. (currently amended) A method of inducing an immune response against Pvs25 on the surface of *Plasmodium vivax* ookinetes, the method comprising administering to a susceptible organism a pharmaceutical composition comprising a nucleic acid encoding a Pvs25 polypeptide in an amount sufficient to induce a transmission blocking immune response, wherein the nucleic acid encodes a Pvs25 polypeptide having at least 95% identity to SEQ ID NO:4.
- 14. (original) The method of claim 16, wherein the susceptible organism is a human.
 - 15-50 (canceled)

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1 51. (new) A composition comprising an isolated nucleic acid molecule which 2 encodes a Pvs25 polypeptide, wherein the polypeptide induces an immune response in a 3 susceptible organism that blocks the transmission of malaria, and wherein the nucleic acid 4 molecule has at least 95% identity to SEQ ID NO:3. 1 52. (new) A composition of comprising an isolated Pvs25 polypeptide having 2 at least 95% identity to SEQ ID NO:4, wherein the polypeptide induces an immune response in a 3 susceptible organism that blocks the transmission of malaria. 1 (new) The composition of claim 52, wherein the Pvs25 polypeptide has 53. 2 an amino acid sequence as shown in SEQ ID NO:4 1 54. (new) A pharmaceutical composition comprising a pharmaceutically 2 acceptable carrier and a Pvs25 polypeptide of claim 52 in an amount sufficient to induce an 3 immune response in a susceptible organism. 1 55. (new) The composition of claim 54, wherein the Pvs25 polypeptide 2 comprises an amino acid sequence encoded by the nucleic acid of claim 1. 1 56. (new) The composition of claim 54, wherein the Pvs25 polypeptide 2 comprises an amino acid sequence encoded by the nucleic acid sequence of SEQ ID NO:3 or an 3 amino acid having the sequence as set forth in SEQ ID NO:4. 1 57. (new) The method of claim 13, wherein the nucleic acid encoding the 2 Pvs25 polypeptide is administered intramuscularly, intradermally, subcutaneously, or 3 intranasally. 1 58. (new) A method of inducing an immune response against Pvs25 on the 2 surface of *Plasmodium vivax* ookinetes, the method comprising administering to a susceptible 3 organism a pharmaceutical composition comprising a Pvs25 polypeptide of claim 52 in an 4 amount sufficient to induce a transmission blocking immune response.

l	59.	(new)	The method of claim 58, wherein the Pvs25 polypeptide in the
2	pharmaceutical comp	osition	is recombinantly produced.
1	60.	(new)	The method of claim 58, wherein the susceptible organism is a
2	human.		
l	61.	(new)	The method of claim 58, wherein the Pvs25 polypeptide in the
2	pharmaceutical comp	osition	is on the surface of a recombinant virus.
l	62.	(new)	The method of claim 58, wherein the Pvs25 polypeptide is
2	administered intramuscularly, intradermally, subcutaneously, or intranasally.		